KATINKO PAIN AND ITCH RELIEVING- camphor (synthetic), menthol, methyl salicylate ointment
Greenstone Pharmaceutical Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Katinko Pain And Itch Relieving Ointment

Active Ingredients:

Camphor 11 percent

Menthol 7.6 percent

Methyl Salicylate 13.0 percent

Purpose

Topical Analgesic

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Uses:

For temporary relief of minor aches and pains of muscles and joints associated with:

Simple Backache Arthritis Rheumatism Muscle Strain Sprains Bruises

For temporary relief of pain and itching associated with:

Insect Bites Minor skin irritation

Warnings

For External Use Only.

When using this product:

Use only as directed.

Do not apply to wounds or damaged skin.

Do not use on the eyes or on the mucous membranes.

Do not use with a heating pad or apply external heat.

Stop use and ask a doctor if:

Skin redness or excessive irritation of the skin occurs

Condition worsens

Symptoms persists for more than 7 days or clears up and occurs again within a few days

Do not bandage tightly

Keep out of reach of children

If swallowed, get medical help or contact a poison control

Directions:

Adults and children 2 yrs and up	Apply to affected areas not more than 3 to 4 times daily
Children under 2 years	Consult a doctor before use

Other Information

This product make provoke allergic reaction in some individuals. Test on small area before use.

Inactive Ingredients

Petroleum Jelly, Paraffin Wax

MANUFACTURED BY:

Greenstone

Pharmaceutical

H.K. Inc.

Anabu Industrial Estate,

Imus, Cavite, Philippines 4103

PRODUCT O F THE PHI L I P PI N E S

MEDICATED

For external use only

30g Package Label

KATINKO PAIN AND ITCH RELIEVING

camphor (synthetic), menthol, methyl salicylate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52241-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	11 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	7.6 g in 100 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
PARAFFIN (UNII: I9O0E3H2ZE)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52241-100-10	1 in 1 CARTON	05/24/2010	
1		10 g in 1 JAR; Type 0: Not a Combination Product		
2	NDC:52241-100-03	1 in 1 CARTON	05/24/2010	
2		3 g in 1 JAR; Type 0: Not a Combination Product		
3	NDC:52241-100-30	1 in 1 CARTON	05/24/2010	
3		30 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/24/2010	

Labeler - Greenstone Pharmaceutical Inc. (719794307)

Registrant - Greenstone Pharmaceutical Inc. (719794307)

Establishment

Name	Address	ID/FEI	Business Operations
Greenstone Pharmaceutical Inc.		719794307	manufacture(52241-100)